

MALARIA

COURSE OBJECTIVES

- BASIC UNDERSTANDING OF MALARIA
 - EPIDEMIOLOGY
 - SYMPTOMS
 - DIAGNOSIS
 - TREATMENT
 - PREVENTION

WHAT IS IT?

- A MOSQUITO-BORNE INFECTIOUS DISEASE OF THE TROPICS, SUBTROPICS, AND FRINGES OF TEMPERATE FORESTS .
 - FOUND IN LATIN AMERICA, CARIBBEAN, ASIA, AFRICA, EUROPE
- CAN BE EITHER ACUTE OR CHRONIC

WHAT IS IT?

- CAUSED BY PROTOZOAN PARASITE GENUS *PLASMODIUM*
- FOUR SPECIES:
 - *P FALCIPARUM*
 - *P VIVAX*
 - *P OVALE*
 - *P MALARIAE*

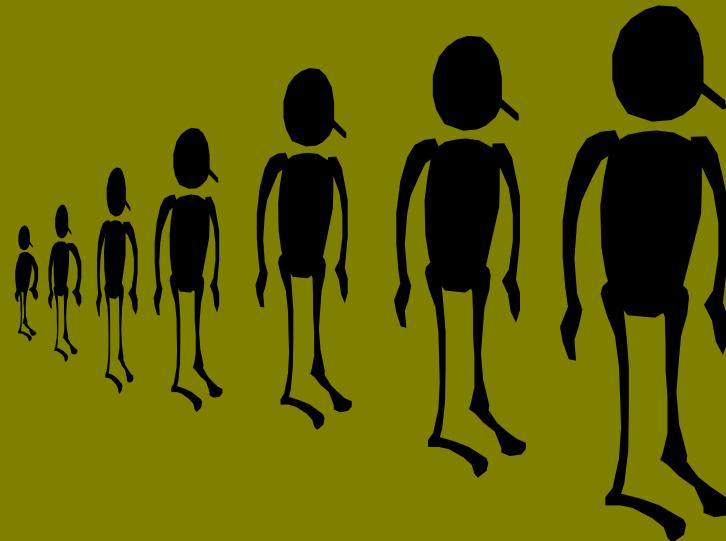
WHY THE CONCERN?

- MOST PREVALENT DISEASE IN THE WORLD
 - 2.1 BILLION LIVE IN MALARIOUS AREAS
 - 100-300 MILLION NEW CASES ANNUALLY
 - 1-3 MILLION DEATHS ANNUALLY
- POTENTIALLY LETHAL DISEASE
- SERIOUS THREAT TO MILITARY OPS



HISTORY

- WWI NAVAL FORCES
 - 4,746 NEW HOSPITAL ADMISSIONS
 - 68,373 LOST MAN-DAYS
 - 7 DEATHS



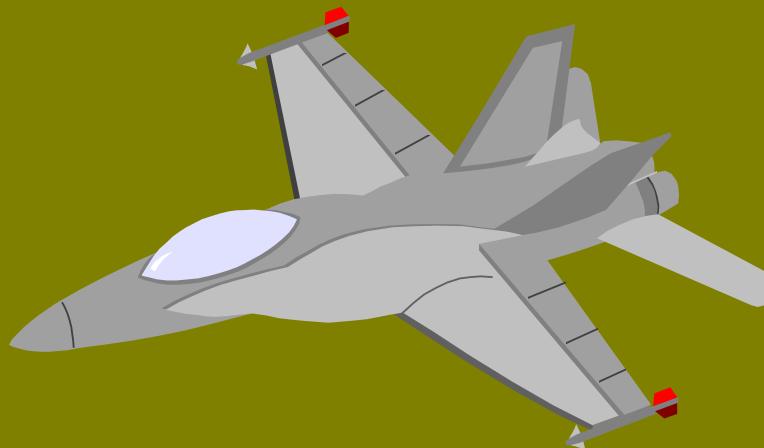
HISTORY

- WWII NAVAL FORCES
 - 111,675 NEW HOSPITAL ADMISSIONS
 - 3,310,800 LOST MAN-DAYS
 - 90 DEATHS
 - 5,332 (AVG) DAILY SICK LIST IN PACIFIC



HISTORY

- VIETNAM WAR
 - 21,695 NEW ADMISSIONS
 - 187,478 LOST MAN-DAYS
 - 46 DEATHS



TRANSMISSION

- MAN IS THE ONLY IMPORTANT RESERVOIR
- VECTOR IS FEMALE ANOPHELES MOSQUITO
 - TEMPERATURE
 - RAINFALL
 - ALTITUDE
 - TERRAIN

TRANSMISSION

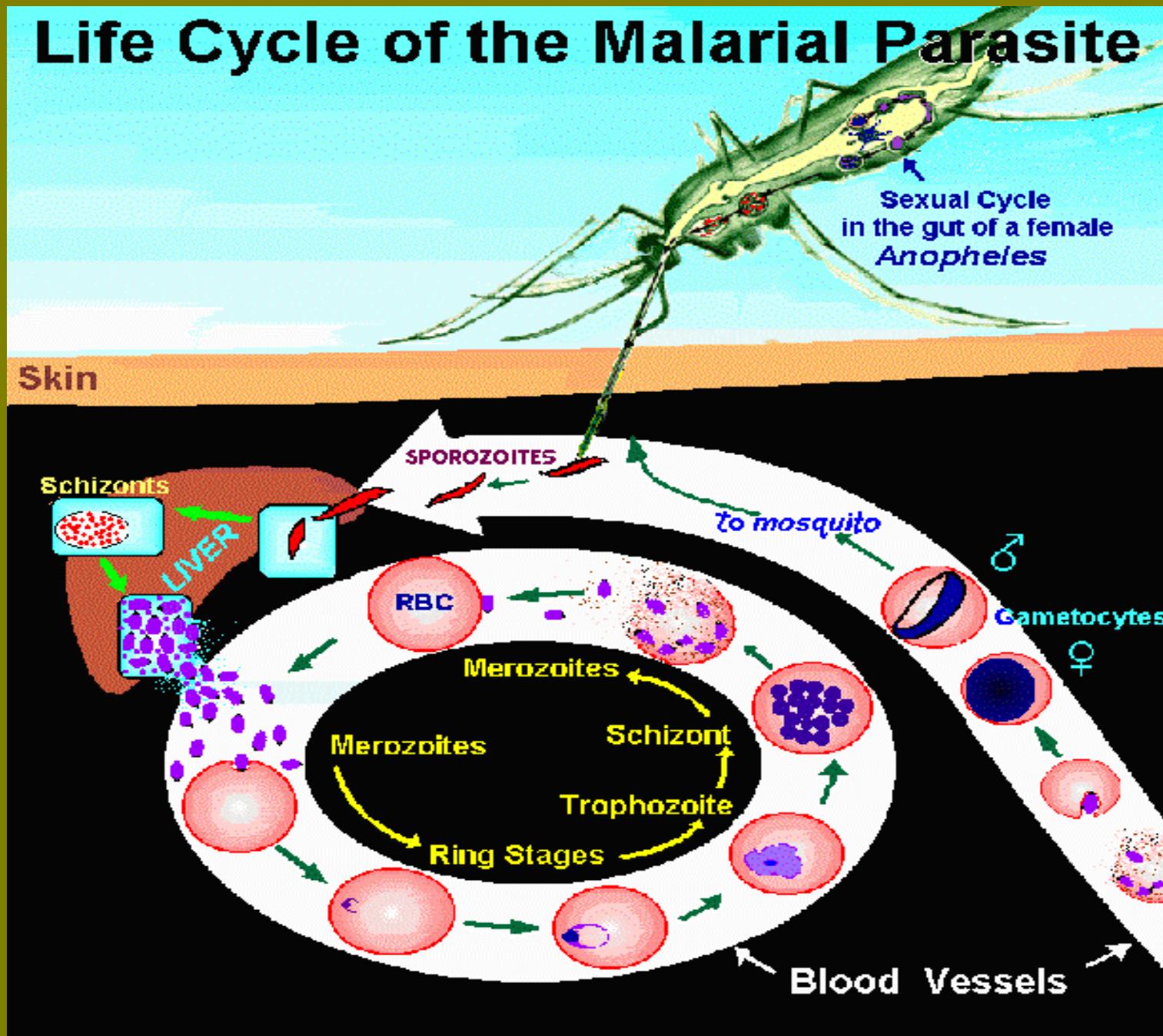
- MOSQUITO VECTOR:
ANOPHELES
- BLOOD TRANSFUSION
- CONTAMINATED NEEDLE
- ORGAN TRANSFER
- CONGENITAL



SUSCEPTIBILITY

- UNIVERSAL SUSCEPTIBILITY
- NO ABSOLUTE IMMUNITY
 - PARTIAL IMMUNITY IN AREAS OF HIGH ENDEMICITY

Life Cycle of the Malarial Parasite



PATHOGENESIS

- RED BLOOD CELL DESTRUCTION
- IMMUNE COMPLEXES AND MEDIATORS
- CAPILLARY PERMEABILITY
- TISSUE HYPOXIA



PLASMODIUM SPECIES

- *P. FALCIPARUM*
 - MOST SEVERE AND PREVALENT
 - 40-60% OF CASES
 - WIDESPREAD CHLOROQUINE RESISTANCE
 - INFECTS RBCs OF ALL AGES: HEAVY PARASITEMIA

PLASMODIUM SPECIES

- P. VIVAX
 - 30-40% OF CASES
 - LIVER PHASE
 - INFECTS YOUNG RBCs: LESS SEVERE THAN FALCIPARUM
- OVALE
 - LIVER PHASE
 - INFECTS YOUNG RBCs
- MALARIAE
 - CAN PERSIST SUBCLINICALLY FOR EXTENDED PERIODS OF TIME
 - INFECTS OLD RBCs

INCUBATION PERIOD

- *P. FALCIPARUM* 12 DAYS
- *P. VIVAX* 14 DAYS*
- *P. OVALE* 14 DAYS*
- *P. MALARIAE* 30 DAYS

* MAY BE 8 - 10 MONTHS OR
LONGER FOR SOME STRAINS

ACUTE SYMPTOMS

- CLASSICAL CYCLIC PAROXYSM:
 - COLD STAGE: CHILLS AND SHAKING
 - HOT STAGE: WARM, HEADACHE, VOMITING
 - SWEATING STAGE: WEAKNESS
 - FEEL WELL FOR PERIOD OF TIME, THEN CYCLE REPEATS ITSELF

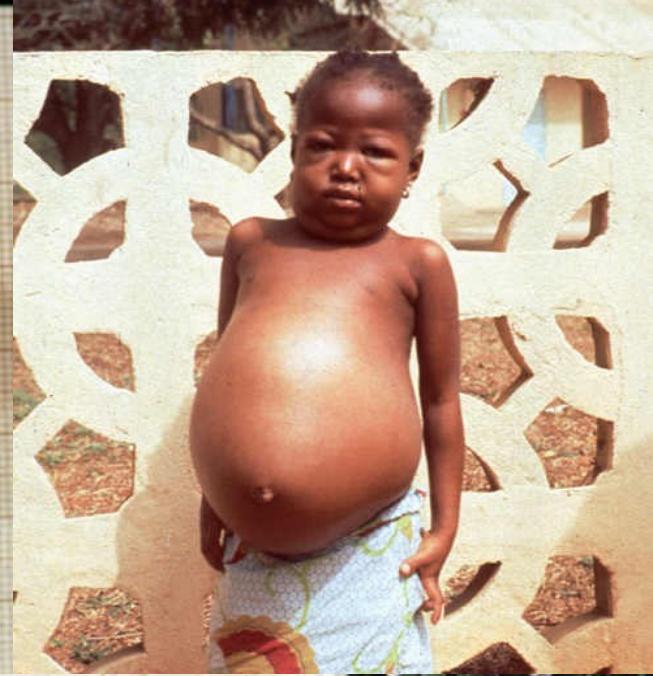
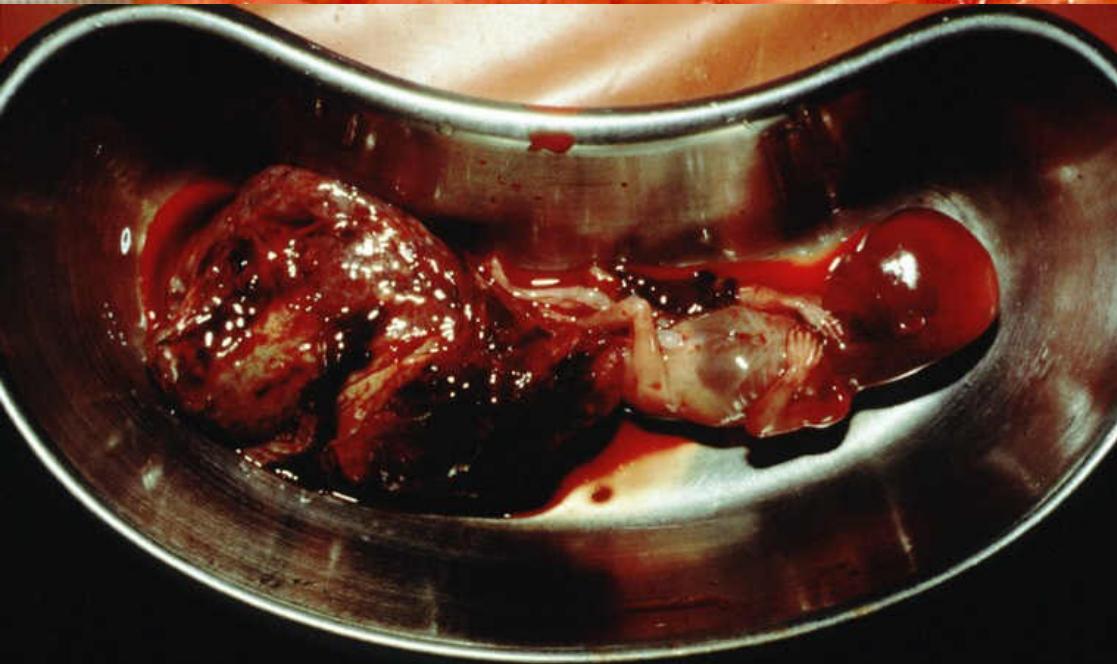
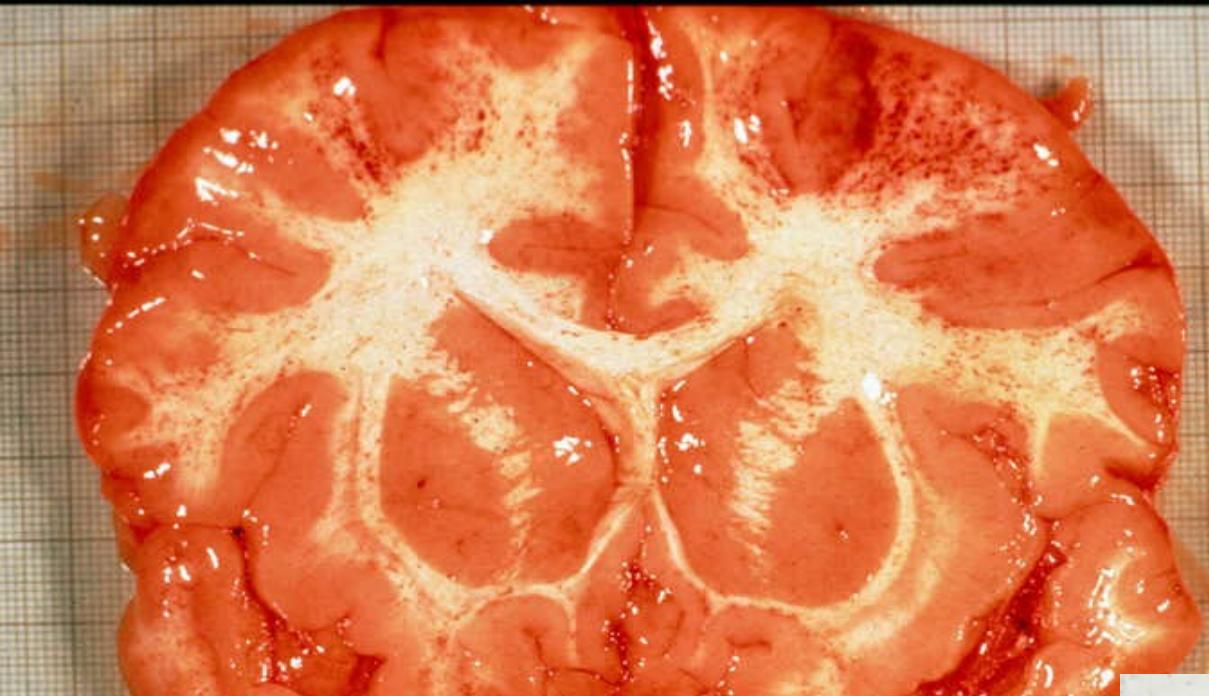
PRESENTATION

- FEVER 97%
- CHILLS 97%
- HEADACHE 97%
- NAUSEA OR VOMITING 62%
- ABDOMINAL PAIN 56%
- MYALGIA 50%
- BACKACHE 9%
- DARK URINE 3%

COMPLICATED MALARIA

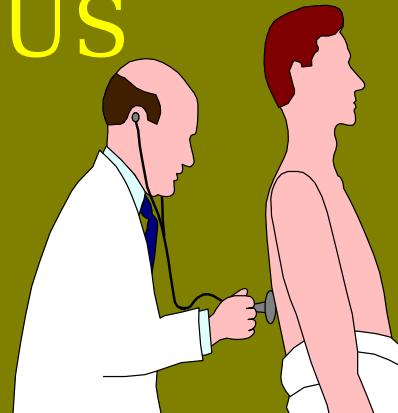
- HYPERPARISITEMIA: >3%
- HYPOGLYCEMIA: <60 MG/DL
- SEVERE ANEMIA:
 - HCT < 21% OR RAPIDLY FALLING HCT
- RENAL FAILURE
- HYPONATREMIA
- CEREBRAL MALARIA
- PROLONGED HYPOTHERMIA
- HIGH OUTPUT VOMITING OR DIARRHEA
- PREGNANCY





SIGNS IN ACUTE INFECTION

- SLIGHTLY ILL TO IN DISTRESS
- ALERT TO UNCONSCIOUS
- FEVER



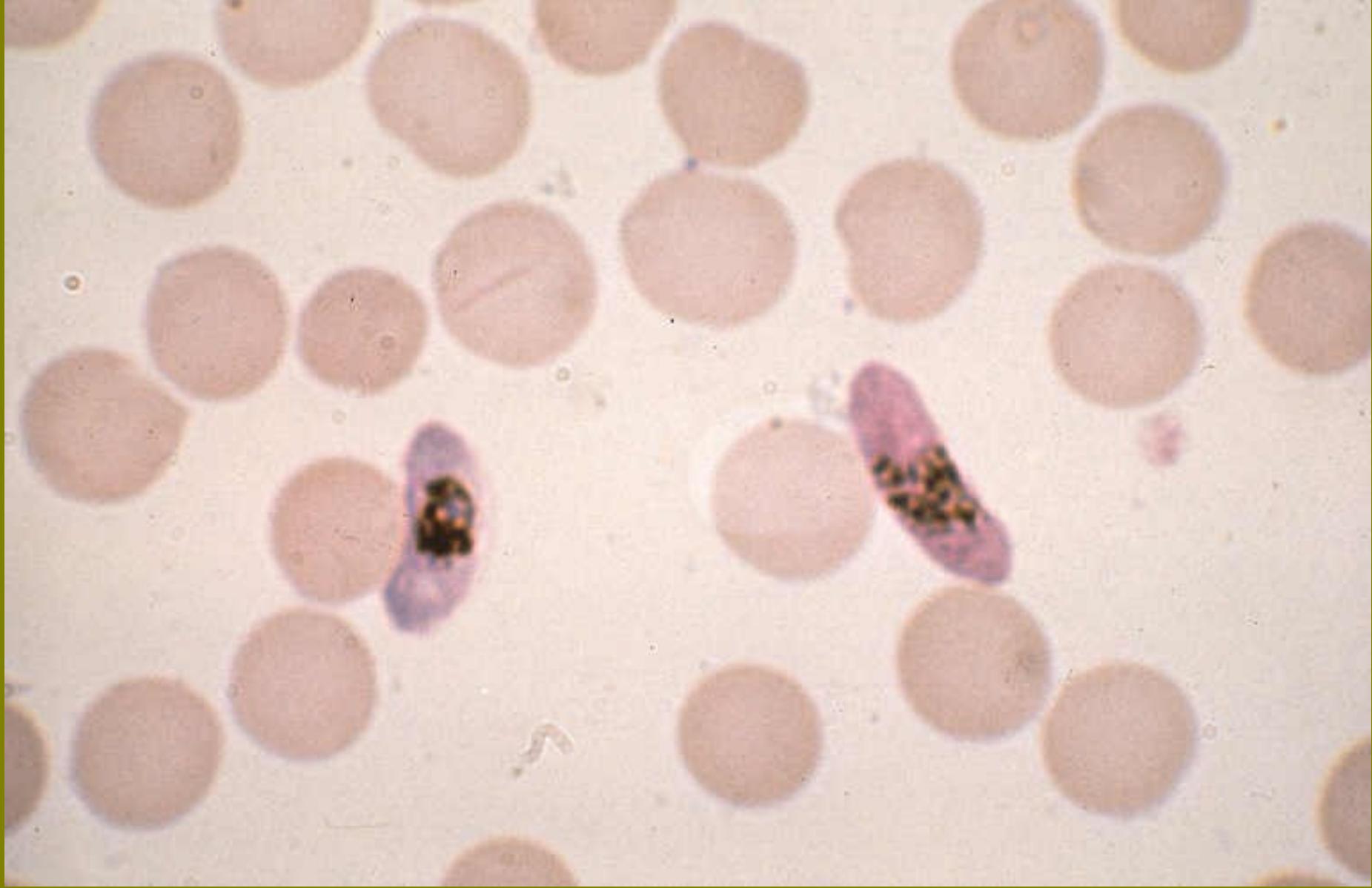
DIAGNOSIS

- **GOLD STANDARD: MULTIPLE THICK AND THIN SMEARS**
- **DIP STICK TESTS**
- **CBC**
 - **ANEMIA**
 - **LEUKOPENIA, OR LEUKOCYTOSIS**
 - **NO EOSINOPHILIA**



Multiple RBC's with trophozoite stage of infection





Schizont stage of malarial infection

TREATMENT

- CHLOROQUINE SENSITIVE INFECTIONS:
 - CHLOROQUINE BASE 600 MG (2 TABS) P.O. INITIALLY, THEN 300 MG (1 TAB) IN 6 HRS, AND QD FOR 2 DAYS
PLUS
 - PRIMAQUINE BASE 30 MG (2 TABS) P.O. PER DAY FOR 14 DAYS

TREATMENT

- UNCOMPLICATED CHLOROQUINE RESISTANT INFECTIONS:
 - QUININE 650 MG PO TID X 3 DAYS AND DOXYCYCLINE 100 MG PO BID X 7 DAYS
- COMPLICATED OR SEVERE INFECTIONS:
 - Intravenous antimalarial medications

TREATMENT

- VIVAX AND OVALE THERAPY
SHOULD INCLUDE PRIMAQUINE
30 MG BASE PO QD X 14 DAYS

OPTIMAL TREATMENT APPROACH

- RAPID CASE IDENTIFICATION
- RAPID PARASITOLOGICAL CLASSIFICATION
- RAPID INITIATION OF THERAPY
- RAPID INITIATION OF SUPPORTIVE CARE

CONTROL OF MALARIA

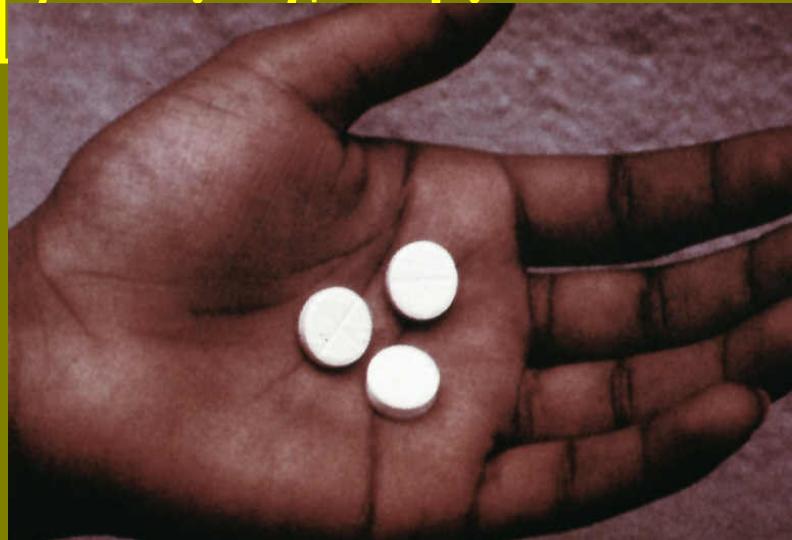
- GLOBAL ERADICATION EFFORTS BY WHO IN 1950s
- EFFORTS NOW FOCUS ON CONTROL vs ERADICATION

POINTS OF ATTACK

1. ATTACK THE PARASITE IN THE HUMAN HOST
2. REDUCE CONTACT BETWEEN HUMANS AND MOSQUITOES
3. DECREASE MOSQUITO POPULATION

ATTACK THE PARASITE IN THE HUMAN HOST

- Treat malaria infections with effective medications
- Use prophylactic drugs to prevent illness and transmission



ATTACK THE PARASITE IN THE HUMAN HOST

- CHEMOPROPHYLAXIS BASED ON CURRENT DRUG RESISTANCE PATTERNS
- **MEFLOQUINE** FIRST LINE PROPHYLAXIS
 - Mefloquine 250mg. po q week, 2 wks prior to exposure and for 4 wks after exposure
- **DOXYCYCLINE** AS SECOND LINE DRUG
 - Doxy 100mg. po qd, 2days prior to exposure and for 4 wks after exposure
- **Primaquine** 30 mg. po qd x 14 days terminal prophylaxis



REDUCE CONTACT BETWEEN HUMANS AND MOSQUITOES

- PERSONAL PROTECTIVE MEASURES

- PROPER WEARING OF UNIFORM
- DEET LOTION
- PERMETHRIN TREATED UNIFORMS
- BED NETS



DECREASE MOSQUITO POPULATION

- ◆ SURVEILLANCE OF MOSQUITO POPULATIONS
- ◆ ID AND ELIMINATION OF BREEDING SITES
- ◆ PROPER INSECTICIDE APPLICATION
 - ATTACK LARVAL STAGES
 - ATTACK ADULT MOSQUITO



SOMALIA

- ♦ **48 CASES IN COUNTRY**
- ♦ **83 CASES FOLLOWING RETURN**
 - 62 USA, 21 USMC



SOMALIA

- ISSUES
 - COMMAND RESPONSIBILITY
 - COMPLIANCE: SWITCH TO MEFLOQUINE
 - LOCATION OF CAMPS
 - WEATHER → POOR USAGE OF DEET
 - NO USE OF BED NETS
 - PRIMAQUINE TERMINAL PROPHYLAXIS
 - USA HAD NOT RECOMMENDED

SOMALIA CONCLUSIONS

- ♦**CASES RESULTED FROM FAILURE TO IMPLEMENT PROPER PROPHYLAXIS AND PERSONAL PROTECTION**
- ♦**MEFLOQUINE MORE EFFECTIVE THAN DOXYCYCLINE IN USMC**



COMMAND RESPONSIBILITY

- Field Marshall Sir William Slim:

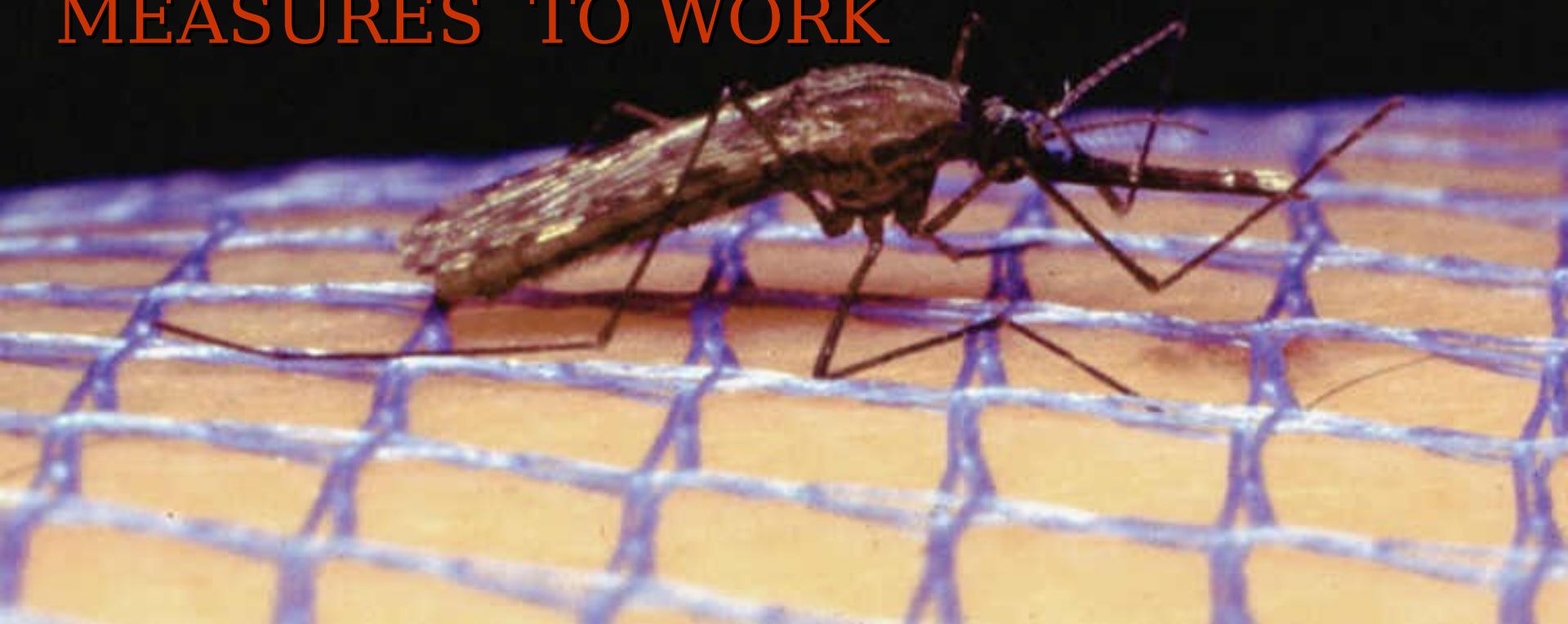
“If the overall result was less than 95% positive, I sacked the commanding officer. I only had to sack three; by then the rest had got my meaning.”
- Sir Neil Cantlie, Dir Gen Brit Army Med Services

“When for the first time in history a combatant officer was considered unfit to command a unit on the grounds that he had allowed his men to become ineffective through disease, a new day in military medicine dawned. The clouds of forgetfulness must not be allowed to overshadow the brightness of that day.”



CONCLUSIONS

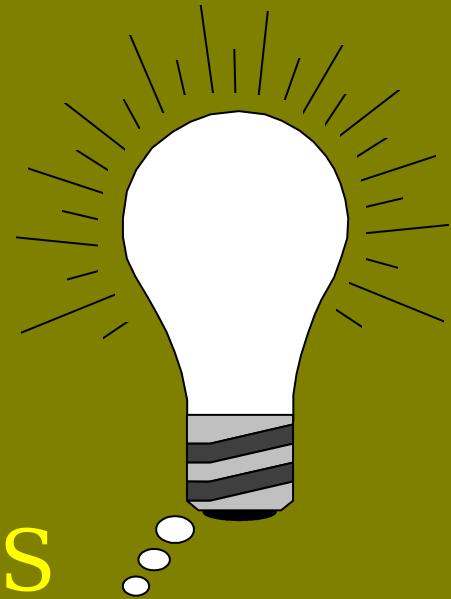
EDUCATION AND AGGRESSIVE
MONITORING OF COMPLIANCE
NEEDED FOR CHEMOPROPHYLAXIS
AND PERSONAL PROTECTIVE
MEASURES TO WORK



SUMMARY

- MALARIA DRUG RESISTANCE IS INCREASING
- GREAT IMPORTANCE OF **PERSONAL PROTECTIVE MEASURES**
- AGGRESSIVE MONITORING NEEDED TO ENFORCE PPM AT COMMAND LEVEL
- DX REQUIRES HIGH INDEX OF SUSPICION
- REGARD AND MANAGE MALARIA AS MEDICAL EMERGENCY

MALARIA!



- REMEMBER:

FLU-LIKE SYMPTOMS

+

'RECENT' HX TRAVEL
TO MALARIOUS AREA

=

THINK MALARIA